S-sulfhydration as a cellular redox regulation

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Synopsis

For many years reactive oxygen and nitrogen species (ROS and RNS) have been recognized as key messengers in the process of thiol-based redox regulation. Relatively recently, literature reports began to mention reactive sulfur species (RSS) and their role in thiol regulation.

This review is focused on biogenesis and biological properties of RSS, including: hydropersulfides, polysulfides and hydrogen sulfide (H$_2$S). Based on the most up-to-date literature data, the paper presents biological significance of S-sulfhydration process. In this reaction, sulfane sulfur is transferred to the -SH groups forming hydropersulfides. Protein cysteine residues, called “redox switches” are susceptible to such reversible modifications. In line with the most recent reports, it was emphasized that sulfane sulfur-containing compounds (mainly hydrogen persulfides and polysulfides) are real and better mediators of S-sulfhydration-based signaling than H$_2$S.

We also overviewed proteins participating in the formation and transport of reactive sulfur species and in mitochondrial H$_2$S oxidation. In addition, we reviewed many reports about proteins unrelated to sulfur metabolism which are modified by S-sulfhydration that influences their catalytic activity. We also addressed the problem of the regulatory function of S-sulfhydration reaction in the activation of K$_{ATP}$ channels (vasorelaxant) and transcription factors (e.g. NF$_{κ}$B) as well as in the mechanism of therapeutic action of garlic-derived sulfur compounds. Some aspects of comparison between RNS and RSS are also discussed in this review.

Key words: hydrogen sulfide; hydropersulfides; reactive sulfur species; S-sulfhydration; sulfane sulfur, sulfurtransferases

Abbreviations:
ROS, reactive oxygen species; RNS, reactive nitrogen species; RSS, reactive sulfur species; GSH, glutathione; Trx, thioredoxin; RSSH, hydropersulfides; CSE, cystathionine ɣ-lyase; MST, 3-mercaptopyruvate sulfurtransferase; CBS, cystathionine-β-synthase; GR, glutathione reductase; SQR, sulfide quinone oxidoreductase; ETHE1, persulfide dioxygenase, Rhd, rhodanese; CAT, cysteine aminotransferase; CDO, cysteine dioxygenase; DAO, D-amino acid oxidase; GAPDH, glyceraldehyde-3-phosphate dehydrogenase; PTP1B, protein tyrosine phosphatase; PTEN, lipid phosphatase; SO, sulfite oxidase; DHLA, dihydrolipoic acid
1. Introduction

Reactive oxygen species (ROS), life threatening products of incomplete oxygen reduction, which damage biomolecules and impair their biological action, are an inevitable consequence of the appearance of oxygen and obligate aerobic organisms on the Earth billions of years ago. Thus, aerobic organisms had to develop a perfect antioxidant defense but simultaneously, they accomplished something more astonishing, namely, they used ROS at low concentrations as regulators of a variety of biological processes. Currently, it is known that the physiological level of oxidants performs regulatory function while their high levels cause oxidative stress and cell damage which leads to pathological states. The commonly accepted key messengers of redox signal transduction in the cell include not only reactive oxygen species (ROS) [1,2] but also reactive nitrogen species (RNS) [3] and reactive sulfur species (RSS) [4,5], which are the principal subject of this review.

1.1. Reactive sulfur species

The concept of reactive sulfur species (RSS) has been postulated in 2001 by Giles et al. [6]. These authors proposed RSS as a group of redox active molecules, which are formed in vivo under conditions of oxidative stress and can act as aggressive oxidizing agents [6]. According to this concept, RSS which are produced under oxidative stress, include thyl radicals (RS•), sulfenic acids (RSOH), disulfides (RSSR), thiosulfinate (RS(O)SR), thiosulfonate (RS(O)2SR) and S-nitrosothiols (SNT). More recently, this definition was expanded to include sulfur-containing molecules, which are formed in physiological, non-oxidative conditions [7,8]. This hypothesis includes also another class of RSS, i.e. the products of cysteine transformations: hydrogen sulfide (H2S) and sulfane sulfur-containing compounds. In the literature, RSS created under physiological conditions (without oxidative stress) are called "the first class of RSS" while "the second class of RSS" means species formed upon the initial action of oxidative stress [7]. However, considering chronology of their appearance, reverse names would be more adequate. In this review, we will discuss processes mediated by RSS created during cysteine transformation, mainly H2S and products of its oxidation – inorganic polysulfides (H2Sn) as well as hydropersulfides (RSSH).

Cysteine sulfur can be metabolized via two main pathways: anaerobic route leading to reduced reactive sulfur species (sulfane sulfur-containing compounds and hydrogen sulfide) and aerobic route yielding sulfates and taurine (Figure 1). In rats, almost one third of the total cysteine sulfur is catabolized to the reduced sulfur compounds while two thirds are oxidized via cysteinyl sulfinate to sulfates and taurine [9]. At present, it is not known which mechanisms underlie the vital decision in the cell about the rate of either the former or the latter process in cysteine metabolism.

1.1.1. Sulfane sulfur – definition, formation and importance

Sulfane sulfur-containing compounds contain a reactive sulfur atom occurring in 0 and -1 oxidation state and bound with another sulfur atom. The following examples of biologically active sulfane-sulfur-containing compounds are worth mentioning: hydropersulfides (RSSH) and hydrogen persulfide (HSSH), thiosulfate (S2O32-), organic (RS,R) as well as inorganic (H2Sn) polysulfides (n≥3) and elemental sulfur (S8) (Figure 1). Sulfane sulfur shows high activity in biological systems while protein carriers which stabilize and transport it are widespread [10,11]. Initially, it was thought that sulfane sulfur plays only a role in
detoxification of cyanide originating from cyanogenic plants (forming thiocyanate) catalyzed by rhodanese, however, this view was substantially changed over the course of time. According to current knowledge, the sulfane sulfur-containing compounds, in particular hydropersulfides (RSSH) and polysulfides (H$_2$S$_n$ and RS$_n$R) are very important for signaling, redox homeostasis, regulation of metabolism and antioxidant defense in the cell [10-12]. Besides the beneficial effects of high activity of some RSS, especially organic polysulfides, their highest concentrations can exert toxic effects that are connected with their ability to generate ROS [13].

Highly metabolically reactive sulfane sulfur-containing compounds are endogenous metabolites created in mammalian cells during anaerobic cysteine catabolic pathway in the reactions catalyzed by such enzymes as cystathionine γ-lyase (CSE), 3-mercaptoppyruvate sulfurtransferase (MST) as well as cysteine aminotransferase (CAT). The anaerobic pathway of L-cysteine conversion can be initiated by CAT-catalyzed transamination of L-cysteine to 3-mercaptoppyruvate (Figure 2). The latter compound is produced also in some tissues from D-cysteine by D-amino acid oxidase (DAO) [14]. 3-Mercaptoppyruvate can play the role of a sulfur donor for different nucleofilic acceptors, including protein sulfhydryl groups with hydropersulfide formation. The sulfur atom is transported from 3-mercaptoppyruvate by 3-mercaptoppyruvate sulfurtransferase (MST) to form MST hydropersulfide (MST-SSH). The level of hydropersulfides, which contain sulfane sulfur, in the cells showing MST and CAT expression is twice higher compared to the cells lacking these enzymes [15]. Moreover, recent studies of Kimura et al. suggest the possibility of inorganic polysulfide production from 3-mercaptoppyruvate. These studies performed in the mouse brain revealed the presence of H$_2$S$_3$ in the cytosol of brain cells and identified MST as the enzyme responsible for its production [16]. The experiments reported in that paper were performed without reducing agents, which could reduce H$_2$S$_n$ to produce H$_2$S. The presence of an acceptor or reducing agent (i.e. mercaptoethanol) can remove H$_2$S from the active site of MST before H$_2$S$_3$ formation is complete. On the other hand, production of H$_2$S$_3$ was strongly suppressed in the presence of a high concentration of substrate (2 mM 3-mercaptoppyruvate), because in the absence of reducing agents, excess of H$_2$S$_3$ in the active site of MST may suppress the progress of reaction, what was shown previously [17]. H$_2$S$_3$ identified by Kimura et al. was also produced from H$_2$S by MST and Rhd [16].

Another pathway of sulfane sulfur formation is associated with cystine conversion into cysteine persulfide called thiocysteine (Cys-SSH) by pyridoxal phosphate (PLP)-dependent cystathionine γ-lyase (CSE) and cystathionine β-synthase (CBS) (Figure 2). The production of thiocysteine from cystine was reported first time by Cavallini et al. [18]. Then, Szczepkowski and Wood demonstrated that thiocysteine could be converted to the sulfane sulfur-containing trisulfide, thiocystine which is a substrate for rhodanese (Rhd) [19] (Figure 2). The detectable cystine concentration in cells is relatively low compared to the extracellular space. This suggests that in the cytosol where CSE and CBS are active, cystine is quickly transformed into thiocysteine. Thiocysteine concentration in cells was estimated at 1-4 μM [5].

It can be concluded that CSE and MST take part in sulfane sulfur formation and transport. In turn, Rhd only transports reactive sulfur atom from donors (i.e. hydropersulfides, trisulfides) to acceptors (other sulfhydryl groups, cyanide). Not only enzymes involved in sulfur metabolism, like CSE, MST, Rhd but also proteins unrelated to sulfur metabolism, such as plasma albumin have the ability to bind and transport sulfane sulfur (Figure 2) [10].
1.1.2. Hydrogen sulfide as a gasotransmitter

The interest in RSS has arisen when hydrogen sulfide (H\textsubscript{2}S) has emerged as a signaling molecule. It happened much later after the signaling role of nitric oxide (NO) and carbon monoxide (CO) had been revealed. Most probably it was related to the conviction that H\textsubscript{2}S was a strong toxin which was upheld till the first reports of the physiological role of this gaseous molecule [20]. Since then reports on diverse biological activities of H\textsubscript{2}S have exploded. Hydrogen sulfide was shown to influence blood pressure, cell proliferation and apoptosis, angiogenesis, inflammatory processes, it was found to be cardioprotective, neuromodulatory and protective against hypoxia, which was described in many review articles [21-24]. These multidirectional physiological actions of H\textsubscript{2}S from the beginning have suggested its implication in diverse signaling processes. However, in spite of those numerous reports highlighting its multitidinous physiological effects, the mechanisms of its actions still have not been fully elucidated.

H\textsubscript{2}S is produced mainly from L-cysteine, however, recently a novel pathway of its production from D-cysteine catalyzed by D-amino acid oxidase has been discovered [14]. The enzymes involved in H\textsubscript{2}S production include PLP-dependent CSE and cystathionine \(\beta\)-synthase (CBS) as well as PLP-independent MST (Figure 2). The expression of the genes for these enzymes is tissue specific and CSE is the major H\textsubscript{2}S-producing enzyme in the cardiovascular system, liver and kidney [22]. H\textsubscript{2}S is formed in the reaction catalyzed by CBS or CSE or non-enzymatically by reduction of hydropersulfides (Figure 2). MST produces H\textsubscript{2}S from 3-mercaptoppyruvate, which is generated from L- or D-cysteine. The sulfur atom is transported from 3-mercaptoppyruvate by MST in the form of MST hydropersulfide (MST-SSH). It was shown that H\textsubscript{2}S could be released in the presence of dithiols (dihydrolipoic acid DHLA, dithioretol, thioredoxin Trx), however, in the presence of monothiols (GSH, cysteine) no release of H\textsubscript{2}S was observed [25]. It could probably be caused by sulfane sulfur transfer from MST-SSH to cysteine or GSH with the formation of Cys-SSH or GSSH. Moreover, sulfur atom of 3-mercaptoppyruvate can be transferred to MST forming not only MST-SSH, but also higher forms of protein polysulfide MST-SS\textsubscript{n}H, from which H\textsubscript{2}S is released in the presence of reducing agents. MST, like Rhd, may also catalyze the reaction of H\textsubscript{2}S production from thiosulfate.

H\textsubscript{2}S differs from other gasotransmitters in its ability to dissociate. Its \(pK_a\) is 6.77 and in physiological conditions, i.e. at pH 7.4 in aqueous solutions, 80\% of hydrogen sulfide is dissociated to HS\textsuperscript{-} anion and only ca. 20\% remains undissociated while S\textsuperscript{2-} concentration is extremely low. It is usually assumed that the hydrogen sulfide pool comprises H\textsubscript{2}S, HS\textsuperscript{-} and S\textsuperscript{2-}. The concentration of H\textsubscript{2}S in human plasma and tissues is still controversial. Initially, reports indicated a high concentration of H\textsubscript{2}S was observed [25]. It could probably be caused by sulfane sulfur transfer from MST-SSH to cysteine or GSH with the formation of Cys-SSH or GSSH. Moreover, sulfur atom of 3-mercaptoppyruvate can be transferred to MST forming not only MST-SSH, but also higher forms of protein polysulfide MST-SS\textsubscript{n}H, from which H\textsubscript{2}S is released in the presence of reducing agents. MST, like Rhd, may also catalyze the reaction of H\textsubscript{2}S production from thiosulfate.
determine an actual H$_2$S level, various new methods of H$_2$S detection have been developed. Zhang et al. reported 0.5 µM concentration of free H$_2$S in human plasma using naphthalimide-azide method with fluorescence detection [31]. Similar free H$_2$S concentrations (below 1 µM) were obtained using the monobromobimane method coupled with RP-HPLC by other authors, who also assayed different forms of sulfane sulfur-containing compounds [32]. The obtained results clearly indicate that the free H$_2$S level in biological systems is much lower that the level of sulfane sulfur containing inorganic polysulfides. It seems that other RSS occurring in tissues at higher concentrations than H$_2$S were responsible for the observed biological effects. It was confirmed by studies in which H$_2$S$_n$ present in the brain activated the transient receptor potential ankyrin 1 (TRPA1) channels approximately 300 times more potently than H$_2$S did [33]. In the light of these findings, small amounts of H$_2$S$_n$ even in the presence of excess of H$_2$S can serve as important biological regulators.

Toohey has indicated that H$_2$S is rather a biodegradation byproduct of sulfane-sulfur-containing compounds [34]. S-sulfhydration reaction with H$_2$S or H$_2$S$_n$ participation plays a crucial regulatory role and at present researches’ interest is focused on the biogenesis of RSS from endogenous and exogenous precursors and on regulatory properties of this reaction.

1.2. Thiol-based redox regulation

Thiol-based redox regulation utilizes a characteristic feature of some redox-sensitive cysteine -SH groups in proteins, called “switches” or “sensors”, which have a unique ability of causing versatile, reversible oxidation to disulfides, sulfenic acids, S-nitrosothiols, mixed disulfides with proteins and hydropersulfides [35-38]. Protein -SH groups can also conjugate with aldehydes forming semiacetals, and can react with different electrophilic groups and metal cations. This capability of diverse modifications of protein -SH groups helps to grasp the mechanism and significance of -SH groups in signaling processes. However, it should be noted that not all protein -SH groups are comparably sensitive to such modification. Sensitivity of different protein cysteine -SH groups to oxidation depends mainly on the pK$_a$ value, i.e. on the ability of its dissociation to thiolate anion (-S⁻), since thiolates are much stronger nucleophiles than thiols. In turn, thiol ionization is facilitated by local environment (positively charged neighboring amino acids, hydrogen bonding and N-terminal α helix). Moreover, some proteins were observed to possess the specialized active-site architecture surrounding cysteine residue that activates substrate binding. For example, H-bonding interactions in peroxiredoxins, besides lowering pK$_{as}$, activate the substrate [39-40]. On the other hand some studies revealed that besides low pK$_{as}$, there were other factors influencing the thiol reactivity.

Kinetic studies reviewed and discussed by Winterbourn and Hampton distinctly indicated that in many cases protein thiols exhibited huge differences in reactivity to an oxidant despite similar pK$_a$ values [37]. Numerous studies demonstrated that only some thiol proteins were oxidized under given conditions and that sensitivity to oxidation depended on an oxidizing agent. Moreover, the compartment where the oxidants (ROS, RNS or RSS) are generated is an important factor of response to oxidative stress due to the possibility of diffusion across the membrane and the distance of the sensitive -SH groups [37-38]. The selected redox-sensitive thiol proteins (sensors, switches), once oxidized, can facilitate the oxidation of other proteins which are implicated in the functional regulation of cell response to redox potential-related environmental changes [36]. When redox-active protein cysteine groups are localized in the active center of key regulatory enzymes, then the redox modulation can control whole metabolic pathways. Some protein -SH groups can undergo reversible oxidation leading to regulatory allosteric effects.
Protein -SH groups can be oxidized not only by one-electron oxidants, i.e. radical species, like ROS, RNS and RSS but also by two-electron oxidants which can show a regulatory effect already at physiological concentrations, that was specifically emphasized by Jones [41-42]. These reversibly oxidized –SH groups are under control of glutathione (GSH), cysteine (Cys) and thioredoxin (Trx), called “nodes” and play an important role in signaling processes. The main function of the “nodes” in redox regulation is to reduce the reversibly oxidized protein thiols, i.e. disulfides, sulfenic acids and S-nitrosothiols back to thiols. Trx is believed to be the main system reducing sulfenic acid while glutaredoxin (Grx) reduces mixed disulfides with proteins [42].

2. Biogenesis and biological properties of hydropersulfides, S-sulfhydration products

Since the time when hydrogen sulfide (H2S) has been recognized to be a gasotransmitter, studies have concentrated not only on biological properties of ROS and RNS but also on RSS generated during anaerobic cysteine metabolism. Currently, the greatest interest in this group is placed on hydropersulfides (RSSH and HSSH) and related inorganic and organic polysulfides (RSnR and H2Sn, n≥3), and H2S, due to their regulatory functions. Metabolism of sulfur amino acids leading to the formation of the above RSS is presented in Figure 2.

It appears that biological actions initially attributed to H2S are actually caused by sulfane sulfur-containing compounds. The formation of RSSH, in which sulfur is in -1 oxidation state, is one of possible reactions leading to reversible oxidation of thiol -SH groups (in which sulfur is in -2 oxidation state). Thus, protein S-sulfhydraltion to hydropersulfides is the oxidation reaction and is believed to be a fundamental process in thiol-based redox regulation which is confirmed by the constantly increasing number of reports of proteins regulated by this process [43-47]. In this review, we focus on thorough characterization of the protein S-sulfhydration reactions and on the significance of this covalent modification of -SH groups in thiol-based redox regulation. Hydropersulfides first drew attention of researchers when they were noticed to act as intermediates in the process of sulfur introduction during biosynthesis of vitamins, cofactors, thionucleosides and iron-sulfur proteins [48]. It indicates that apart from the regulatory function, cysteine-derived sulfane sulfur also participates in biosynthesis of key molecules in many biochemical processes.

It is worth mentioning that there are significant differences between properties of -SH and -SSH groups, namely, hydropersulfides (-SSH) show much greater chemical reactivity compared with the corresponding thiols (-SH) due to the increased nucleophilicity. At physiological pH, the acid-base balance of hydropersulfides in shifted towards persulfide anion which means that hydropersulfides are stronger acids (lower pKₐ value) and better hydrogen donors than their thiol counterparts [43]. Another significant difference is associated with much lower dissociation energy of the S-H bond in RSSH which equals 70 kcal/mol while the corresponding value for the S-H bond in RSH amounts to 92 kcal/mol [49]. It can be attributed to the increased stability of perthiyl radicals (RSS•) which are stabilized by resonance compared to thiyyl radicals (RSS•). All these features make hydropersulfides very efficient antioxidants. For instance, a remarkable increase in the reducing capacity of GSSH relative to GSH was observed [5]. Toohey attributes this fact to the possibility of GSSH tautomerization to form thiosulfoxide, which easily donates electrons [50] (reaction 1).

\[
S \rightleftharpoons 2 \text{GSSH} \rightarrow 2 \text{GSH} \rightarrow 2e^- + 2H^+ + \text{GSS-SSG} \quad (1)
\]
A recent study by Ida et al. confirmed that only GSSH (generated from GSSSG by glutathione reductase) could scavenge a marked amount of H$_2$O$_2$, while comparable amounts of GSH (formed in the reaction of glutathione reductase with GSSG) produced no measurable H$_2$O$_2$ reduction [5]. It confirms that more nucleophilic hydropersulfides are superior reducing agents compared with their thiol counterparts. Moreover, these findings show that the sulfane sulfur in hydropersulfides is an extremely reactive reductant and is a potent antioxidant in cells. These authors underline that compared with GSH and H$_2$S, hydropersulfides (mainly GSSH and CysSSH as well as protein hydropersulfides) are more efficient nucleophiles and reductants which fulfill critical regulatory functions in redox signaling in the cell [5].

Hydropersulfides are unstable compounds and in the presence of thiols can undergo desulfhydration accompanied by H$_2$S release or disproportionation reaction yielding thiol and elemental sulfur S$_8$ (designated also as S$^0$) [51] (reaction 2-3).

\[
\text{RSSH} \xrightarrow{R'SH} \text{RSSR'} + \text{H}_2\text{S} \quad (2)
\]

\[
\text{RSSH} \rightarrow \text{RSH} + \text{S}^0 \quad (3)
\]

For this reason, sulfhydration can be also perceived as a storage method for the toxic H$_2$S and as a path for elemental sulfur (S$_8$) release [52]. The fact that H$_2$S is released from hydropersulfides in the presence of thiols explains why initially many biological functions were attributed to H$_2$S. Numerous data from experimental studies indicate that rather not H$_2$S but other reactive sulfur metabolites, mainly hydropersulfides or inorganic polysulfides, are capable of producing antioxidant and regulatory effects. What is more, it is increasingly more often emphasized that biological activity of H$_2$S is mostly connected with the formation of hydropersulfides. Ida [5] and Toohey [34] even believe that this is not H$_2$S but sulfane sulfur in a wider sense that fulfills an essential role in thiol-based redox regulation. However, many studies continue to report biological effects of S-sulfhydration by the H$_2$S. Probably it results from the fact that, as mentioned earlier, all commercial reagents (Na$_2$S or NaHS) contain significant amounts of sulfane sulfur. Results of Greiner et al. revealed the inhibition of PTEN activity by reversible oxidation of this enzyme by all tested H$_2$S donors (i.e. Na$_2$S or NaHS from various companies as well as the polysulfide K$_2$S$_x$), and demonstrated that the degree of inhibition depended on the sulfane sulfur content [53]. The use of the “purest” preparation resulted only in a slight lowering of the activity (by less than 10%), while commercial preparations containing a considerable amount of sulfane sulfur caused even 60% inhibition [53].

Quantitative data indicate that hydropersulfides and inorganic polysulfides are widespread in cells and tissues and occur at much higher physiological concentrations than ROS or RNS. This confirms a crucial role of S-sulfhydration reaction in regulatory processes [5,45], and this is why this process and its biological significance attract increasing interest.

2.1. Hydropersulfide formation

The hydropersulfides (RSSH) are formed by the oxidation of thiols (from -2 to -1 oxidation state). It is surprising, as also noticed by Greiner [53], that many reports do not take notice of the fact that sulfur in H$_2$S and in HS$^-$ can act only as a reductant, and its direct participation in oxidation reactions of -SH groups to hydropersulfides (RSSH) is impossible. Therefore, hydropersulfides cannot be formed in a direct reaction of H$_2$S with protein -SH groups. Hence, the S-sulfhydration reaction can occur only when one of the reagents (-SH group or H$_2$S) is in the oxidized form.
The cysteine group can be S-sulfhydrated by HS\textsuperscript{−}, when its -SH group is oxidized to sulfenic acid, disulfide, mixed disulfide or nitrosothiol \cite{54-55}. The S-sulfhydration process of proteins consisting in the nucleophilic attack of HS\textsuperscript{−} anion on the reversibly oxidized -SH groups is presented in reactions 4-7. In these reactions, the sulfur of hydrosulfide anion is transformed into sulfane sulfur present in the generated hydropersulfide.

\[
\begin{align*}
\text{Protein-SOH} + \text{HS}^- & \rightarrow \text{Protein-SS}^- + \text{H}_2\text{O} \quad (4) \\
\text{Protein-SNO} + \text{HS}^- & \rightarrow \text{Protein-SS}^- + \text{HNO} \quad (5) \\
\text{Protein-S} + \text{HS}^- & \rightarrow \text{Protein-SS}^- + \text{SH} \quad (6) \\
\text{Protein-SSG} + \text{HS}^- & \rightarrow \text{Protein-SS}^- + \text{CS}^- \quad (7)
\end{align*}
\]

It appears that in cytosolic environment which is reducing due to the glutathione system, the protein -SH groups are rarely oxidized, thus this mechanism of the sulfhydration will not be preferred. However, when GSH is deficient or in more oxidizing environment, like mitochondria or endoplasmic reticulum those reactions can occur. The formation of hydropersulfide of sulfide quinone oxidoreductase (SQR) during mitochondrial H\textsubscript{2}S oxidation is an example of the S-sulfhydration reaction via the above mechanism (see 2.3). Recently, Vasas et. al. performed kinetic studies on disulfide reduction reaction by HS\textsuperscript{−} \cite{56}. Those authors demonstrated that the reaction occurred via multistep equilibria through disulfide, hydropersulfide and inorganic polysulfides (reaction 8-10). They suggested that reduction of disulfide (RSSR) by HS\textsuperscript{−} was a highly system-specific process, and thermodynamic and kinetic premises indicated a tight regulation. Those authors proposed that chemical nature of the disulfide moiety would play a major role in the feasibility of its reduction by hydrosulfide anion \cite{56}. Although those studies were conducted on non-protein disulfides (DTNB, CysSSCys, GSSG), it appears that the results can be applied to protein disulfides. In proteins, the reaction will depend on the surrounding amino acids. The presented mechanism indicates that inorganic polysulfides (HS\textsubscript{n}−) are formed in parallel with hydropersulfides (RSS\textsuperscript{−}).

\[
\begin{align*}
\text{RSSR} + \text{HS}^- & \rightarrow \text{RSS}^- + \text{RSH} \quad (8) \\
\text{RSS}^- + \text{HS}^- & \rightarrow \text{RS}^- + \text{HSS}^- \quad (9) \\
\text{nHSS}^- + \text{HS}^- & \rightarrow \text{HS}_{(n+1)}^- + (n-1) \text{HS}^- \quad (10)
\end{align*}
\]

Recently, Cuevasanta et al. in their excellent work studied the reactivity of HS\textsuperscript{−} toward symmetric disulfides (RSSR) and mixed disulfides with albumin (HSA). They observed the twentyfold increase in HSA persulfide reactivity in comparison to the HSA thiol at pH 7.4 \cite{57}. It shows a very improved nucleophilicity of persulfides with respect to the parent thiols and H\textsubscript{2}S. Experiments with cells in culture indicate that hydropersulfide formation increases
upon exposure to hydrogen peroxide. These authors presented also kinetic evidence that the hydropersulfides have much better nucleophilic reactivity due to alpha effect [57].

S-sulfhydration reaction can also occur via reaction of protein -SH group with oxidized forms of hydrogen sulfide (HSOH or HSSH) (reaction 11-12).

\[
\begin{align*}
\text{HSOH} & \quad \text{Protein-S}^- + \text{H}_2\text{O} \quad \text{Protein-SS}^- \quad (11) \\
\text{HSS}^- & \quad \text{Protein-S}^- + \text{HS}^- \quad \text{Protein-SS}^- \quad (12)
\end{align*}
\]

Nagy and Winterbourn investigated the reaction of HS\(^-\) with hypochlorous acid (HOCl). This reaction leads initially to sulfenyl chloride which undergoes hydrolysis yielding sulfenic acid and then condensation with another HS\(^-\) anion to produce polyhydropersulfides [58] (reaction 13). Products of this reaction can be oxidizers of protein -SH groups which leads to the formation of protein hydropersulfide (reaction 12).

\[
\begin{align*}
\text{HS}^- \xrightarrow{\text{HOCI}} & \quad \text{HS-Cl} \xrightarrow{\text{H}_2\text{O}} \quad \text{HS-OH} \xrightarrow{\text{HS}^- \quad (\text{HS}^-)_{n}} \quad \text{H(S)\textsubscript{n}S}^- \quad n=1-7 \quad (13)
\end{align*}
\]

Moreover, as already mentioned, H\(_2\)S can be easily oxidized to inorganic polysulfides, so sulfide and hydrosulfide salts are usually contaminated with sulfane sulfur. Thus, inorganic polysulfides can be intermediates in the -SH to -SSH transformation pathway and participate in the thiol-based redox signaling [53,59]. Sulfane sulfur present in polysulfides can be easily transferred to thiolate anion with hydropersulfide formation [59]. So, when we add sulfide solution to a protein solution, in fact we introduce the sulfane sulfur. Based on the above observations, it is not difficult to imagine that indeed inorganic polysulfides and hydropersulfides can be responsible for biological activity of H\(_2\)S. Moreover, some of RSS containing sulfane sulfur (mainly hydropersulfide and thiosulfate) are generated during mitochondrial sulfide oxidation pathways (see 2.3) and in this way they can modify activity of target proteins.

Hydropersulfides can also be produced via the radical pathway [29]. One electron oxidation of H\(_2\)S leads to HS\(^-\) formation, which can next react with protein thiols to give finally hydropersulfide and superoxide radical anion in accordance with reactions 14-15.

\[
\begin{align*}
\text{HS}^- + \text{Protein-SH} & \longrightarrow \text{Protein-SSH}^- + \text{H}^+ \quad (14) \\
\text{Protein-SSH}^- + \text{O}_2 & \longrightarrow \text{Protein-SSH} + \text{O}_2^- \quad (15)
\end{align*}
\]

2.2. Involvement of thiocysteine in the glutathione and protein S-sulfhydration process

Protein S-sulfhydration can occur also as the result of sulfane sulfur transfer from low molecular weight hydropersulfides to protein -SH groups. Thiocysteine, generated from
cystine by CSE or CBS (reaction 16), can be involved in this S-sulfhydration process (Figure 2).

\[
\text{CSE, CBS} \quad \begin{array}{c}
\text{Cys-S-S-Cys} \\
\rightarrow \\
\text{Cys-SS}^\text{H} + \text{pyruvate} + \text{NH}_3
\end{array} \quad (16)
\]

Moreover, thiocysteine sulfane sulfur can be transferred to glutathione (GSH), present in cells at high concentrations, with the formation of glutathione hydropersulfide (GSSH) (reaction 17) [5].

\[
\text{GSH} \\
\rightarrow \\
\text{Cys-SSH} \quad \rightarrow \\
\text{GSSH} + \text{Cys-SH} \\
\quad (17)
\]

According to literature data, the GSSH concentration in mammalian tissues is high (over 100 μM in the brain and about 10-100 μM in other organs [5,54]), which justifies the conclusion that GSSH is the most widespread low molecular weight hydropersulfide, playing a dominant role in S-sulfhydration reactions. However, it seems that the published physiological hydropersulfide concentrations are only estimates because appropriate analytical standards are lacking due to exceptional instability of those compounds.

Ida et al. [5] demonstrated that GSSH could react with glutathione disulfide (GSSG) which led to the formation of the trisulfide (GSSSG), and then in subsequent reactions even higher glutathione polysulfides could be generated (reaction 18).

\[
\text{GSSH} \\
\rightarrow \\
\text{GSSG} \quad \rightarrow \\
\text{GSSSH} \quad \rightarrow \\
\text{GSSSG} \quad \rightarrow \\
\text{GSSSH} \quad \rightarrow \\
\text{GSH} + \text{Cys-SH} \\
\quad (18)
\]

Those authors revealed that all glutathione polysulfides (GSSG, GSSSG, GSSSSG) could be reduced in the reaction catalyzed by glutathione reductase (GR) leading to the generation of GSH-based forms of hydropersulfides (GSSH, GSSSH, GSSSSH). The high GSSH content is maintained by CSE and CBS generating CysSSH and by GR activity that regenerates GSSH and GS(S)_nH from oxidized glutathione polysulfides in cells [5].

In the next steps, GSSH and other low molecular weight hydropersulfides can transfer sulfane sulfur to other -SH groups, most of all to the -SH groups of proteins [5,45,54]. The reactions leading to the exchange of -SH and -SSH groups, like the reaction of GSSG with protein -SH groups, are called S-transsulfhydration reactions. It indicates that the formation of protein hydropersulfides in cells can be a result of sulfane sulfur transfer from low-molecular-weight compounds (reaction 19) to a specific redox-sensitive cysteine residue in the target protein. Many data indicate that CysSSH and then GSSH are the main sources of sulfane sulfur for S-sulfhydration reactions of proteins despite the fact that there are reports of hydropersulfide formation in the reaction of HS^− anion with reversibly oxidized -SH groups.

\[
\text{GSSH} \\
\rightarrow \\
\text{GSH} + \text{Cys-SH} \\
\quad (19)
\]

Sulfane sulfur present in protein hydropersulfides (Protein-SSH) can then be transferred to acceptor -SH groups of other proteins (transsulfhydration) (reaction 20) [5,54], which represents the thiol-based regulatory mechanism.
The protein hydperoxysulfide formation can prevent the accumulation of dangerous H$_2$S excess, and provides a cellular regulatory mechanism of its level. It is commonly believed that protein hydperoxysulfides are a storage form of H$_2$S, which can be released e.g. under the influence of thiols (reaction 21) (desulphydration process) [60].

\[
\text{Protein'-SH} \quad \xrightarrow{\text{Protein-SSH}} \quad \text{Protein'-SSH + Protein-SH} \quad (20)
\]

\[
\text{RSH} \quad \xrightarrow{\text{Protein-SSH}} \quad \text{Protein-SSR + H}_2\text{S} \quad (21)
\]

2.3. Hydrogen sulfide oxidation pathway: hydperoxysulfides as the intermediates

Since interest in biological actions of hydperoxysulfides and hydrogen sulfide (H$_2$S) is on the rise, it would be fundamental to know not only H$_2$S biosynthesis but also biodegradation paths. Since H$_2$S is an active molecule and in higher concentrations it is very toxic, its cellular level has to be tightly regulated because it is able to suppress the respiratory chain by inhibiting cytochrome oxidase. It means that an imbalance between H$_2$S biogenesis, its storage capacity in the hydperoxysulfide form (-SSH) and biodegradation is dangerous. Thus, maintaining the physiological concentration of H$_2$S in tissues is of crucial importance. H$_2$S is biodegraded mostly in mitochondria through a series of oxidations via hydperoxysulfides, sulfite, thiosulfate and sulfate [61]. Mitochondrial H$_2$S oxidation is catalyzed by the following enzymes: sulfide quinone oxidoreductase (SQR), persulfide dioxygenase (ETHE1) and rhodanese (Rhd) [62].

The H$_2$S degradation pathway begins with the reaction catalyzed by sulfide quinone oxidoreductase (SQR). This enzyme catalyzes a two electron oxidation of H$_2$S to the sulfane sulfur containing persulfide SQR-SSH [63-64]. This reaction is like a gate to the H$_2$S oxidation pathway. In this reaction, ubiquinone is an electron acceptor, and this links H$_2$S catabolism with oxidative phosphorylation and makes H$_2$S the first inorganic substrate for electron transfer chain [65]. This indicates that mitochondrial H$_2$S catabolism is conjugated with ATP biosynthesis. SQR is an inner mitochondrial membrane-bound flavoprotein belonging to the class of disulfide oxidoreductases. This enzyme is characterized by a low $K_m$ value (μM) and a high catalytic sulfide turnover rate, supporting efficient catabolism of the toxic substrate [64]. A physiological acceptor of sulfane sulfur from SQR-SSH has not been identified unequivocally, yet. Some authors have postulated that human SQR utilizes sulfite as a persulfide acceptor, yielding thiosulfate as a product, thus, this reaction is rapid and highly efficient at physiological pH (Figure 3, path 1) [63]. On the other hand, studies of Libiad et al. [61] have demonstrated that in addition to sulfite, glutathione (GSH) functions as a persulfide acceptor for human SQR leading to GSSH (Figure 3, path 2). It is not excluded that there are SQR persulfide acceptors other than sulfite and glutathione (e.g. DHLA, Trx, cysteine), which after accepting sulfane sulfur can be reduced by GSH (Figure 3, path 3 presented for DHLA) [66]. Thus, depending on the route, GSSH is the next direct or indirect metabolite on the H$_2$S oxidation pathway. It is the known substrate of persulfide dioxygenase (ETHE1), the second enzyme participating in sulfide oxidation. ETHE1 is a non-heme iron-containing protein in mitochondrial matrix that catalyzes the oxidation of the persulfide, consuming molecular oxygen [67-68]. Sulfite formed in the ETHE1-catalyzed reaction can be oxidized to sulfate by
sulfite oxidase or can accept persulfide sulfane sulfur yielding thiosulfate in the reaction catalyzed by Rhd (Figure 3). An inherited mutation in persulfide dioxygenase gene results in an autosomal recessive disorder called ethylmalonic encephalopathy (EE). In patients suffering from deficient ETHE1 activity, the level of H$_2$S and S$_2$O$_3^{2-}$ rises while that of SO$_3^{2-}$ declines. Toxic sulfide concentrations cause the vascular endothelium damage and lead to neurological failure, chronic diarrhea and disturbances in mitochondrial energy metabolism [69]. Since the hydropersulfide GSSH is an ETHE1 substrate, this enzyme should be perceived not only as a H$_2$S biodegradation tool but also as an enzyme controlling the hydropersulfide level and thus regulating the degree of protein S-sulfhydration in cells.

As mentioned above, rhodanese is the third crucial protein in the H$_2$S oxidation route, (sulfurtransferase, Rhd). Rhd is a mitochondrial enzyme known for its ability to utilize sulfane sulfur in CN$^-$ detoxification reactions with thiocyanate formation [70]. Another substrate of rhodanese, thiosulfate can be generated, for instance, by sulfane sulfur transfer from SQR-SSH to sulfite. Some authors have proposed that thiosulfate is the main substrate for glutathione-dependent Rhd (thiosulfate sulfurtransferase) [71]. This reaction produces glutathione persulfide and regenerates sulfite. However, studies of Hildebrandt and Grieshaber [62] and Libiad et al. [61] have clearly indicated that Rhd preferentially synthesizes rather than utilizes thiosulfate. The $K_m$ values obtained in those studies have strongly suggested that sulfite and persulfides rather than thiosulfate and cyanide are preferable natural substrates for Rhd.

The main products of H$_2$S metabolism and forms of sulfur excreted from the body include thiosulfate and sulfate. In the liver, sulfate is a predominant end product of H$_2$S metabolism in the presence of GSH, while in the absence of GSH, a mixture of sulfate and thiosulfate is generated. Under conditions of high H$_2$S levels, an elevated level of blood and urinary thiosulfate was observed [72]. Interestingly, the S-sulfhydration reaction occurs twice in the mitochondrial H$_2$S oxidation process, namely during the formation of the hydropersulfides SQR-SSH and GSSH. Sulfane sulfur belongs not only to the structure of hydropersulfides but also to thiosulfate (SSO$_3^{2-}$) created in the Rhd-catalyzed reactions. The whole process of H$_2$S oxidation appears to follow a surprisingly tangled path. However, the seemingly complicated scheme of H$_2$S oxidation is biologically justified. Recently, Mishanina et al. in their very interesting paper posited that sulfide oxidation pathway should be considered not only as a mechanism for utilization of excess sulfide, but principally as the way for generation of reactive sulfur species, including persulfides, polysulfides and thiosulfate [8].

Thiosulfate formation can be viewed as one of the methods of H$_2$S storage in cells, from which it can be released by thiosulfate reductase. The elegant experiments of Koj et al. [73] and Skarżyński et al. [74] support the central role for thiosulfate as a key intermediate in the H$_2$S oxidation. Since, as commonly known, sulfate is the final product of aerobic cysteine sulfur metabolism, the thiosulfate concentration in body fluids and tissues is assumed to be an indicator of H$_2$S production in the body.

H$_2$S catabolism is tightly connected with mitochondrial respiratory chain, thus, it would be interesting to know how this compound is degraded by red blood cells lacking mitochondria. Recent studies of Vitvitsky et al. [75] discovered a novel mechanism of H$_2$S oxidation in erythrocytes. They revealed that RBCs utilized methemoglobin to catalyze H$_2$S oxidation producing thiosulfate and polysulfide (Figure 4). The postulated mechanism explains how erythrocytes maintain low H$_2$S levels in circulation, moreover, it cannot be excluded that additional hemoproteins might be involved in maintaining the H$_2$S homeostasis.
2.4. S-sulfhydration of proteins

Not only enzymes involved in sulfur metabolism, like CSE, MST or Rhd [10-11,76] but also proteins entirely unrelated to sulfur metabolism have the ability to bind and transfer sulfane sulfur in the form of hydropesulfides or trisulfides. They include, for instance, the above-mentioned plasma albumin which transports sulfane sulfur in the form of hydropersulfide [10]. The important role of albumin in sulfane sulfur transport was proven by the presence of sulfane sulfur in organs with no or trace enzymatic activity related to sulfane sulfur biosynthesis [77].

Numerous enzymes have been reported to be S-sulfhydrated in vitro and in intact cells, including actin, glyceraldehyde-3-phosphate dehydrogenase (GAPDH) [44], protein tyrosine phosphatase (PTP1B) [78], lipid phosphatase (PTEN) [53], Ni-containing carbon monoxide dehydrogenase [79] Cu/Zn superoxide dismutase [80], malate dehydrogenase [81] and 5-aminolevulinate synthetase [82].

In these enzymes, -SH groups function as sensors (switches) able to bind sulfane sulfur and thus to regulate their activity by hydropersulfide formation. The S-sulfhydration in the active center is also required for the activation of xanthine oxidase and aldehyde oxidase which was demonstrated already in the 1970s [83-84]. Studies on these enzymes suggested that sulfane sulfur occurring in the form of hydropersulfides was a natural component of these proteins. Removal of sulfane sulfur by cyanide treatment (cyanolysis reaction) inactivated those enzymes (reaction 22). The cyanolysis reaction is a proof of the presence of -SSH group in a protein and is a fundamental characteristic reaction of hydropersulfides and other forms of sulfane sulfur.

\[
\text{Enzyme-SS}^- + \text{CN}^- \rightarrow \text{Enzyme-S}^- + \text{SCN}^- \quad (22)
\]

Correspondingly, the incubation of malate dehydrogenase with thiosulfate and Rhd, i.e. an enzyme transforming -SH into -SSH groups, was shown to activate this enzyme (reaction 23) [81].

\[
\text{Enzyme-S}^- + \text{Na}_2\text{S}_2\text{O}_3 \rightarrow \text{Enzyme-SS}^- + \text{Na}_2\text{SO}_3 \quad (23)
\]

Studies of Mustafa et al. revealed that 39 of the liver proteins are sulfhydrated under physiological conditions including GAPDH, β-tubulin and actin, and S-sulfhydration of these proteins can be reversed by dithiotreitol (DTT). Sulfhydration augments GAPDH activity and enhances actin polymerization [44]. The S-sulfhydration reactions in those experiments were dependent on the H_2S endogenous generation potential or its exogenous supplementation. Other studies showed that reaction of Cu/Zn superoxide dismutase (SOD) with sodium sulfide led to the formation of persulfide group at Cys^{111}. This modification made the acid-induced denaturation of SOD fully irreversible [80]. Ni-containing carbon monoxide dehydrogenases from bacteria catalyze the reversible oxidation of CO to CO_2 in the active site containing nickel, iron and sulfur. Studies of Kim et al. have revealed that the persulfide bond is essential for the stability and catalytic activity of the Ni-Fe-S clusters [79]. The protein tyrosine phosphatase B1 (PTPB1) is a member of the protein tyrosine phosphatase family, which regulates various signal transduction pathways. This enzyme has the Cys residue in the active center that is very susceptible to reversible oxidation. In vitro and in vivo studies demonstrated that PTPB1 could be sulfhydrated in its active site which resulted in the inhibition of phosphatase activity. Inactivation of PTPB1 by H_2S (probably containing sulfane sulfur) was reversed preferentially by thioredoxin [78]. Greiner and coworkers have studied
oxidation of protein thiol groups via S-sulfhydration reaction, using lipid phosphatase (PTEN) as a protein model [53]. The activity of this enzyme strictly depended on the free thiol of Cys\(^{124}\) and oxidative modification of this -SH group led to PTEN inactivation. It was found that the addition of sulfane sulfur to cysteine in the active center of PTEN resulted in the inhibition of the enzyme. When PTEN was treated with H\(_2\)S in the presence of GSH, its inhibitory effect on PTEN activity was significantly weaker because GSH prevented formation of the hydropersulfide form in the PTEN active center. Additionally, recent studies of Ohno et al. revealed that both Cys\(^{71}\) and Cys\(^{124}\) in PTEN were the targets for S-sulfhydration. Further, the CBS knockdown in human neuroblastoma cells SH-SY5Y reduced this modification and made PTEN more sensitive to modification by NO [85].

The above examples showed that S-sulfhydration of enzymes affects their activity, which means that it has a regulatory character. Unlike other kinds of modifications of –SH groups (e.g. S-nitrosation, S-glutathionylation), S-sulfhydration often increases catalytic activity of proteins, however, studies on PTPB1 and PTEN indicated that this modification could also suppress their activity.

In the above-mentioned studies, Greiner et al. evidenced that inorganic polysulfides, formed in NaHS solutions, were the oxidizing species responsible for PTEN S-sulfhydration. All tested “H\(_2\)S donors”, like Na\(_2\)S, gaseous H\(_2\)S and GYY4137, led to polysulfide-mediated oxidation of PTEN via addition of sulfane sulfur atom to an important Cys residue in this protein [53]. Those and other studies clearly suggest that the effects previously attributed to H\(_2\)S are rather mediated by inorganic polysulfides and other compounds containing sulfane sulfur. Actual documented low concentrations of H\(_2\)S in tissues and higher levels of sulfane sulfur-containing compounds additionally confirm that the latter compounds play an essential role in the thiol-based redox regulation. However, other physiological actions of H\(_2\)S cannot be excluded, like its interaction with NO and CO and metal ions, and interactions with hemoproteins, e.g. hemoglobin or cytochrome oxidase c.

2.4.1. Regulatory action of S-sulfhydration in the K\(_{\text{ATP}}\) channel activation process

ATP-sensitive potassium (K\(_{\text{ATP}}\)) channels are multi-subunit protein complexes distributed on the surface of the cell and mitochondrial membranes of many different cell types. K\(_{\text{ATP}}\) channels are octameric complexes of two types of membrane-protein subunits. They are composed of four pore-forming Kir subunits and four regulatory subunits known as SUR (sulfonylurea receptor). Each Kir subunit associates with one SUR subunit [86]. Transmembrane Kir subunits allow for K\(^+\) ion influx into the channel complex, while SUR subunit plays a receptor role for different pharmacological compounds which activate or inhibit the channel opening [86]. A drop in the ATP level and a rise in cellular ADP level are physiological triggers of channel opening while sulfonylurea derivatives are pharmacological K\(_{\text{ATP}}\) channel openers.

It is strongly suggested that K\(_{\text{ATP}}\) channels are one of the major targets of H\(_2\)S and are activated via S-sulfhydration (Figure 5) [87-88]. All studies on the K\(_{\text{ATP}}\) channel S-sulfhydration suggest that this process is triggered by H\(_2\)S, however, it should be remembered that these studies utilized Na\(_2\)S or NaHS, which contain significant amounts of sulfane sulfur. To verify which RSS are actually responsible for the biological effects, it would be necessary to carry out comparative studies between the effect of Na\(_2\)S and analogous amounts of inorganic polysulfides.

The activation of K\(_{\text{ATP}}\) channels in vascular smooth muscle cells involves the formation of hydropersulfide form of the protein, which reduces its ATP binding affinity and results in vasorelaxation [88]. Mustafa et al. evidenced that vasorelaxant action of H\(_2\)S was connected with sulfhydration at Cys\(^{43}\), i.e. one of nine cysteine residues in Kir6.1 subunit [88]. On the
other hand, some authors suggested that \( \text{H}_2\text{S} \) sulfhydrated \( \text{SUR2B} \) subunit, not \( \text{Kir6.1} \) subunit [89-90]. Irrespective of which \( \text{K}_{\text{ATP}} \) channel subunit is the main \( \text{H}_2\text{S} \) target, hydropersulfide formation plays a crucial role in potassium channel activation. Moreover, some studies demonstrated that sulfhydration of \( \text{SUR2B} \) subunit modified tyrosine nitration in \( \text{Kir6.1} \) subunit, what suggests the interaction between these two posttranslational modifications [90].

Inhibition of potassium channel by the specific inhibitor glibenclamide blocks vasorelaxant action of \( \text{H}_2\text{S} \) (or RSS) [88]. Likewise, the application of propargylglycine, an inhibitor of the \( \text{H}_2\text{S} \) and sulfane sulfur synthesizing enzyme in vascular cells (i.e. CSE), blocks vasorelaxant action of \( \text{H}_2\text{S} \). The CSE knockout mice show vascular dysfunction and atherosclerosis which facilitates progression of coronary artery disease [91]. For this reason, the drugs increasing endogenous production of \( \text{H}_2\text{S} \) and sulfane sulfur, like N-acetylcysteine, and \( \text{H}_2\text{S} \) precursors can be beneficial in antihypertensive therapy.

### 2.4.2. S-sulfhydration reaction in the transcription factor NF\( \kappa \)B activation process

Signals from hormones, growth factors, cytokines and neurotransmitters are transduced in cells and transmitted to the transcription machinery in the nucleus by a class of proteins called transcription factors, and NF\( \kappa \)B is their representative. This antiapoptotic factor is inactive under basal conditions because it is bound to the inhibitor I\( \kappa \)B (I\( \kappa \)BNF\( \kappa \)B). The multifunctional proinflammatory cytokine, tumor necrosis factor alpha (TNF\( \alpha \)) activates I\( \kappa \)B kinase which leads to its phosphorylation and degradation and to the release of NF\( \kappa \)B and its translocation to the nucleus (Figure 6). Sen and coworkers showed that TNF\( \alpha \)-stimulated transcription of CSE and led to S-sulfhydration of NF\( \kappa \)B. \( \text{H}_2\text{S} \) generated in CSE-catalyzed reactions creates hydropersulfide at Cys\( ^{38} \) in the p65 subunit, which contributes to transcription of antiapoptotic proteins in the nucleus [92]. Therefore, TNF\( \alpha \)-induced S-sulfhydration of p65 subunit of NF\( \kappa \)B improves survival of cells. Some studies showed a dramatic increase in \( \text{H}_2\text{S} \) production in PLC/PRF/5 hepatoma cells compared to human LO2 hepatocyte cells. Moreover, treatment of these hepatoma cells with NaHS as a \( \text{H}_2\text{S} \) donor, markedly increased CSE and CBS expression leading to NF\( \kappa \)B activation, decreasing the number of apoptotic cells and increasing cell viability [93].

### 2.5. Capability of hydropersulfide, sulfane sulfur and hydrogen sulfide formation from garlic-derived sulfur compounds

Therapeutic potential of garlic has been known for a long time. Garlic is known to counteract atherosclerosis, reduce glucose and cholesterol level, strengthen the immune system, and to show anticancer, antioxidant and hypotensive activities [94-96]. The main organosulfur compound occurring in garlic called allicin is unstable and rapidly decomposes mostly to diallyl disulfide (DADS) and diallyl trisulfide (DATS). Benavides et al. have demonstrated that garlic owes its vasoactive effect to the transformation of diallyl polysulfides to the sulfane sulfur-containing hydropersulfides, which react with thiols to release \( \text{H}_2\text{S} \) [97].

The reaction of DATS, which originally has sulfane sulfur in its structure, with GSH produces the mix disulfide allylglutathione and the low molecular weight hydropersulfide allylperthiol (reaction 24), from which \( \text{H}_2\text{S} \) is released in the reaction with GSH [97] (reaction 25).

\[
\text{CH}_2=\text{CH-CH}_2\text{-S-S-S-CH}_2\text{-CH}=\text{CH}_2 \rightarrow \text{CH}_2=\text{CH-CH}_2\text{-SSG} + \text{CH}_2=\text{CH-CH}_2\text{-SSH} \quad (24)
\]
In turn, the reaction of DADS with GSH yields S-allylglutathione and allylperthiol (reaction 26), which reacts with GSH releasing H$_2$S according to reaction 25 [97].

\[
\text{CH}_2=\text{CH}-\text{CH}_2-\text{SSH} \rightarrow \text{CH}_2=\text{CH}-\text{CH}_2-\text{SSG} + \text{H}_2\text{S} \quad (25)
\]

In the above reactions, protein thiols can compete in cells with GSH and can react with allylperthiol which may lead to covalent modification of proteins and formation of mixed disulfides. Moreover, DATS can directly transfer reactive sulfane sulfur to protein -SH groups which generates protein hydropersulfides. Greiner et. al. showed that DATS efficiently inhibited PTEN activity by oxidation of its thiol group in HEK293T cells [53]. On the other hand, DADS, which is a disulfide, can be a source of sulfane sulfur and can transfer this active sulfur atom in the tautomerization reaction to thiosulfoxide (reaction 27) [10].

\[
\begin{align*}
\text{CH}_2=\text{CH}-\text{CH}_2-\text{SSH} & \xrightarrow{\text{GSH}} \text{G-S-CH}_2=\text{CH}_2 + \text{CH}_2=\text{CH}-\text{CH}_2-\text{SSH} \quad (26)
\end{align*}
\]

Hence, it seems that studies on biological and pharmacological activity of garlic are connected with hydropersulfides, sulfane sulfur and H$_2$S.

3. Interconnections and comparison between activities of reactive nitrogen species (RNS) and reactive sulfur species (RSS)

Studies of recent years unraveled in greater depth the biogenesis and regulatory mechanisms governing ROS- and RNS-based responses to different stimulators. Our knowledge of RSS is much more patchy because these compounds, like H$_2$S, attracted researchers’ interest much later. Investigations of NADPH oxidase or NO synthase isoforms with different specific inhibitors contributed much to a better understanding of the regulatory role of ROS and RNS. They also furthered a detailed analysis of redox-sensitive signaling pathways and identification of target molecules, which opened up novel therapeutic perspectives. For this reason, it is advisable to continue analogical studies of RSS which will allow us to fully understand the role of these messengers of redox signaling. For decades it has been known that S-nitrosation of protein -SH groups to S-nitrosothiols (SNT) is a regulatory mechanism [3,98], like S-thiolation that leads to generation of mixed disulfides with proteins [99]. In contrast, regulatory function of protein S-sulfhydration yielding protein hydropersulfides (PSSH) began to focus more interest only recently. Nevertheless, the studies have fully documented that the formation of protein hydropersulfides is a fundamental process in cellular thiol-based redox regulation. This conviction is confirmed by a still increasing number of reports of proteins modified by S-sulfhydration. The significance of this process is also evidenced by the widespreadness and abundance of endogenous glutathione and protein hydropersulfides [5].
There are many functional similarities and interconnections between NO and H$_2$S. It is known that H$_2$S and NO mutually affect their concentrations [100]. Since hydropersulfides can be also created in reactions of S-nitrosothiols (SNT) with HS$^-$ anion, thus, the generation of protein SNT is an intermediate step leading finally to more stable hydropersulfides occurring in cells at higher concentrations. It indicates that S-nitrosation can play the role of a driving force of the S-sulfhydration reaction and both these oxidative reactions can modify the same cysteine residues in proteins. NO reduces blood pressure via cGMP while vasorelaxant action of H$_2$S (or RSS) is executed via S-sulfhydration of K$_{ATP}$ channels. Intermolecular S-transnitrosylation reactions consist in NO migration between -SH groups of different molecules while transsulfhydration involves an analogical migration of sulfane sulfur. The formation and accumulation of GSSH can serve to transfer sulfane sulfur to the preferred acceptor protein -SH groups (with a low pK$_{a}$ value) in S-sulfhydration reactions. Another similarity between S-sulfhydration and S-nitrosation reactions in mammalian cells is that on the one hand, there are three NOS isoenzymes responsible for NO biosynthesis and on the other, there are three main enzymes responsible for biogenesis of H$_2$S and hydropersulfides, i.e. CSE, CBS and MST. All these enzymes differ in tissue distribution and substrate specificity and all produce hydropersulfides facilitating then S-sulfhydration of protein and non-protein thiols. S-sulfhydration, like S-nitrosation, is a reversible process via the thioredoxin system. Thus, the next similarity is occurrence of denitrosation reaction (e.g. S-nitrosoglutathione (GSNO)) and desulfhydration reaction (e.g. GSSH). The difference is that the SNT concentration in tissues and body fluids is low while the level of hydropersulfides is much higher [5]. Thus, S-sulfhydration seems to be a more common regulatory mechanism than S-nitrosation. Both these processes can be compared to the phosphorylation/dephosphorylation regulatory process.

4. Future directions and prospects

Although the current knowledge of the role of S-sulfhydration process in the regulatory mechanisms constantly expands, there are still many problems which await explanation. First of all, it is necessary to explain whether RSS or H$_2$S (or maybe both) are the actual signaling molecules. It should be emphasized that it is more and more often suggested that the sulfane sulfur-containing compounds and not H$_2$S, are real mediators of S-sulfhydration-based signaling. Hence, future studies should make a distinction between biological effects caused by H$_2$S and by sulfane sulfur, and the search for new exogenous, safe sulfane sulfur and H$_2$S donors with pharmacological potential should be continued.

Moreover, it would also be interesting to elucidate to what extent the biological effect can be attributed to HS$^-$ anion and to what extent to H$_2$S. H$_2$S is known to easily cross cell membrane [101] while it was considered to be impermeable for HS$^-$ anion. However, the recent report showed a HS$^-$-permeable channel in bacterial cells [102]. Then, studies of Jennings using the human erythrocyte membrane revealed HS$^-$ influx in exchange for Cl$^-$, catalyzed by the anion exchange protein AE1. These results showed that H$_2$S and HS$^-$ can mediate the transport of acid equivalents across the biological membrane analogically to the cycle for Cl$^-$, HCO$_3^-$ and CO$_2$ [103].

Secondly, the identification of target molecules able to undergo S-sulfhydration should be continued, which may pave the way for novel prophylactic and therapeutic interventions. Some problems related to specificity of S-sulfhydration reactions, vital in signaling processes, also requires further explanation. Specificity studies require precise estimations of the levels of individual RSS in different subcellular compartments. Thus, development of more precise,
sensitive and easy-to-use methods for determination of RSS would be essential for the measurement of their local levels in specific locations within the cell.

The next problem awaiting full elucidation is related to the interaction between H$_2$S and NO, and maybe also with CO. All available evidences indicate that these three gasotransmitters cannot be treated as separate molecules but as cooperating factors involved in physiological function of cells in health and disease [104-105]. Therefore, further studies should explain how the three gasotransmitters cooperate to influence the biosynthesis and activity of different reactive species.

Nonetheless, it is worth underlining that the last decade has witnessed a tremendous advance in our knowledge of the relevance of S-sulfhydration reaction [2,4-5]. It has been evidenced that this process elicits an extraordinarily widespread influence on different biochemical events and biological processes. Therefore, it can be expected that the next decade will bring a better understanding of the intriguing world of chemistry and biochemistry of reactive sulfur species. There are all indications that further progress in this field and new discoveries and new research methods already can be seen on the horizon.

Most recently, Mishanina et al. in their excellent work proposed a new, interesting concept of RSS biogenesis. They suggested that H$_2$S oxidation pathway, considered mainly as a way for disposing the toxic H$_2$S excess, should be regarded as the generation route for RSS that could modify target proteins [8]. In fact, as mentioned earlier, mainly hydropersulfides of proteins (SQRSSH) or low-molecular thiols (GSSH) are formed during H$_2$S oxidation. GSSH can transfer its sulfane sulfur to target protein as shown in Figure 3. Moreover, thiosulfate, the final product of H$_2$S oxidation, is a stable compound containing sulfane sulfur which can transfer this reactive sulfur onto other -SH groups in Rhd-catalyzed reaction. Sulfite produced during H$_2$S oxidation is also proposed to be RSS due to the possibility of sulfite radical anion (SO$_3^{•-}$) formation. Sulfite oxidation to sulfate or its utilization by Rhd can be the way of limiting the damaging potential of sulfite. Authors compared H$_2$S oxidation pathway to the mitochondrial electron transfer, underlining that both are the source of reactive species, namely RSS and ROS, respectively. Moreover, H$_2$S oxidation taking place in erythrocytes catalyzed by methemoglobin (Figure 4) also yields RSS (thiosulfate and metal-bound hydropersulfide) [75]. So, this very interesting hypothesis gives a new look at the importance of H$_2$S oxidation pathway and RSS signaling potential.

**References:**

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Figure Legends

Figure 1. Aerobic and anaerobic transformation of L-cysteine. Reactive sulfur species (RSS) formed in physiological, non-oxidative conditions are marked in gray boxes (S*- sulfane sulfur atom). CDO – cysteine dioxygenase; CAT – cysteine aminotransferase; CSE – cystathionine γ-lyase; MST – 3-mercaptopropylsulfate sulfurtransferase; CBS – cystathionine β-synthase.

Figure 2. Generation and transport of sulfane sulfur and H2S production from L- and D- cysteine in the reactions catalyzed by cysteine aminotransferase (CAT), 3-mercaptopropylsulfate sulfurtransferase (MST), cystathionine-γ-lyase (CSE), cystathionine β-synthase (CBS), rhodanese (Rhd) and D-amino acid oxidase (DAO). Sulfane sulfur-containing compounds are marked in bold. GSH – glutathione, LA – lipoic acid, DHLA – dihydrolipoic acid, Trx – thioredoxin.

Figure 3. Formation of hydropersulfides during mitochondrial H2S oxidation. Thiosulfate and sulfane are the main products of H2S oxidation. Enzymes participating in this process include: SQR – sulfide quinone oxidoreductase, ETHE1 – persulfide dioxygenase, Rhd – rhodanese, SO – sulfite oxidase, DHLA – dihydrolipoic acid.

Figure 4. Scheme of MetHb-dependent H2S oxidation to thiosulfate and inorganic polysulfides bound to heme iron proposed by Vitvitsky et al. ([70] modified).

Figure 5. S-sulfhydration of KATP channel leading to its activation and to vasodilatation. RSS – reactive sulfur species.

Figure 6. S-sulfhydration of the transcription factor NFκB and antiapoptotic gene transcription (according Paul et al. [37] modified). CSE - cystathionine-γ-lyase, RSS – reactive sulfur species.
L-cysteine

**aerobic pathway**

- CDO
- CAT

**anaerobic pathway**

- CSE
- MST
- CBS

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**cysteinesulfinate**

- $+H_3N-CH_2$
- $CH_2$
- $SO_3^-$

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**sulfane sulfur**

- $H_2S_n/H\hat{S}_n^-$
- inorganic polysulfide
- $R-S-\hat{S}H/\hat{RSS}^-$
- hydropersulfide
- $\hat{S}=SO_3^{2-}$
- thiosulfate
- $R-S-\hat{S}_n^-S-R$
- organic polysulfide
- $\hat{S}_8$
- elemental sulfur

---

**hydrogen sulfide**

- $H_2S/HS^-/S^{2-}$
- hydrogen sulfide

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RSS (H₂S) → S-sulfhydration

K⁺ channel

Hyperpolarization → Vasodilation → Blood pressure
TNFα receptor

activation of IkB kinase

IκB

IkB fosforylation and degradation

S-sulfhydration

NFκB

CSE translation

CSE transcription

H2S (RSS)

antiapoptotic gene transcription

Cys38-SSH

IkB

p65

p50

Nuclear membrane

Cell membrane

Accepted Manuscript

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